

Docket: 7214.08

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

First Named Inventor:	Charles N. Serhan	
Appln. No.:	Filed herewith	
Filed:	October 19, 2001	Previous Examiner: Dwayne C. Jones
Title:	Regulation of Phospholipase D Activity	Previous 1614 Group Art Unit:

PRELIMINARY AMENDMENT

Commissioner for Patents
BOX PATENT APPLICATION
Washington, D.C. 20231

Sir:

Please preliminarily amend the above-identified application as follows:

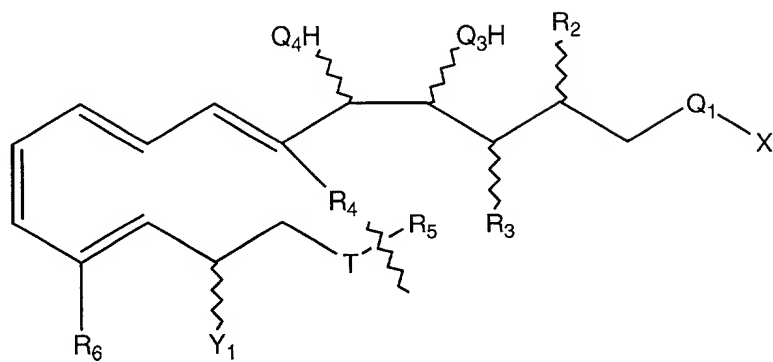
In the specification:

At page 1, lines 3 through 5, please delete the entire paragraph after “Cross-Reference to Related Applications” and replace with the following paragraph:

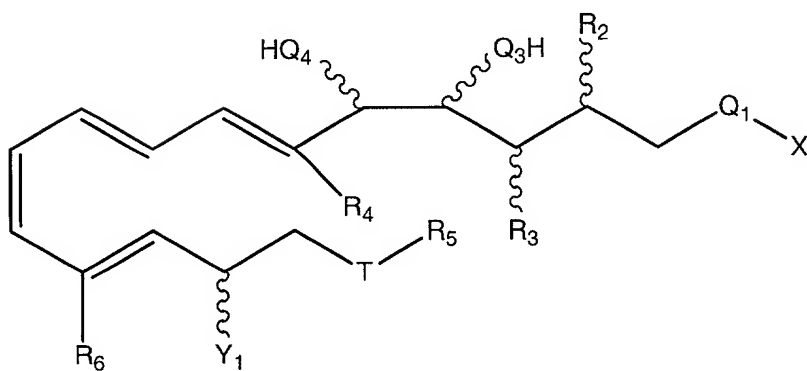
- - This application is a continuation application of U.S. Patent Application No. 09/525,157, filed March 14, 2000, which in turn claims priority to U.S. Provisional Patent Application No. 60/125,194, filed March 18, 1999, the contents of which are incorporated herein by reference. - -

Patent Application No. 09/525,157

At page 11, lines 5 through 10, please delete

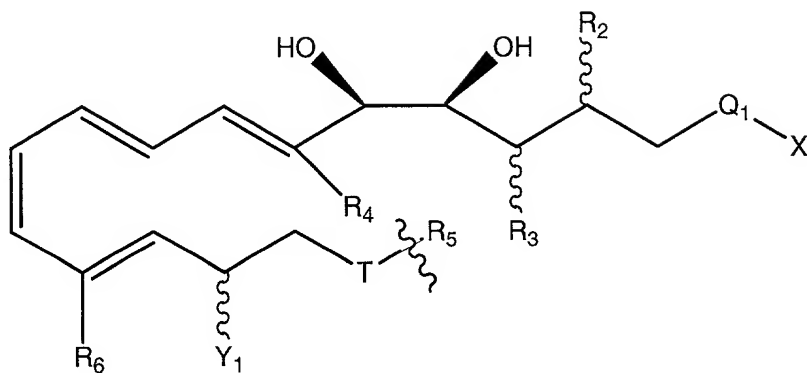


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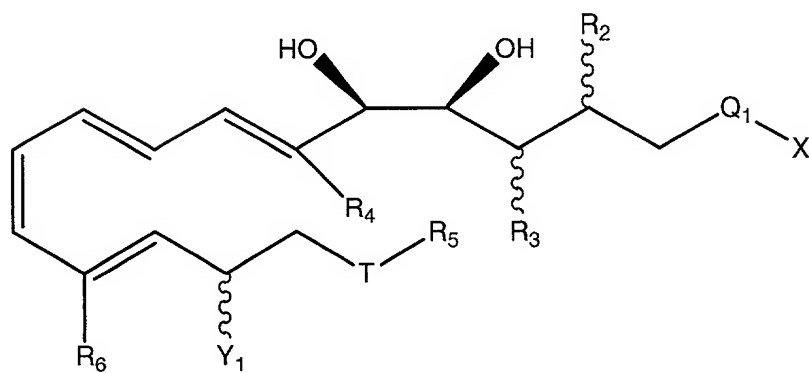


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At page 14, lines 1 through 7, please delete

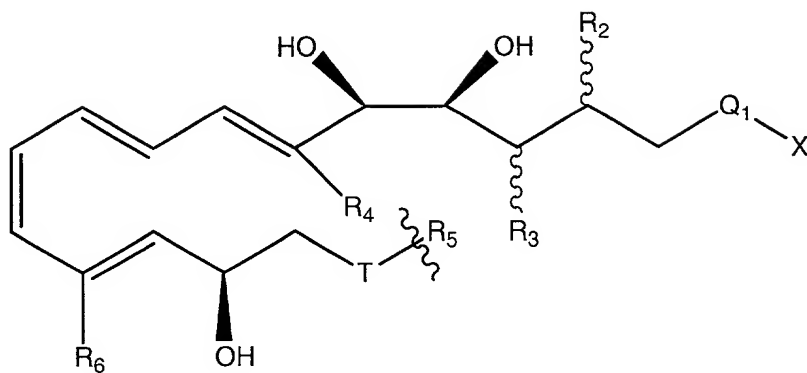


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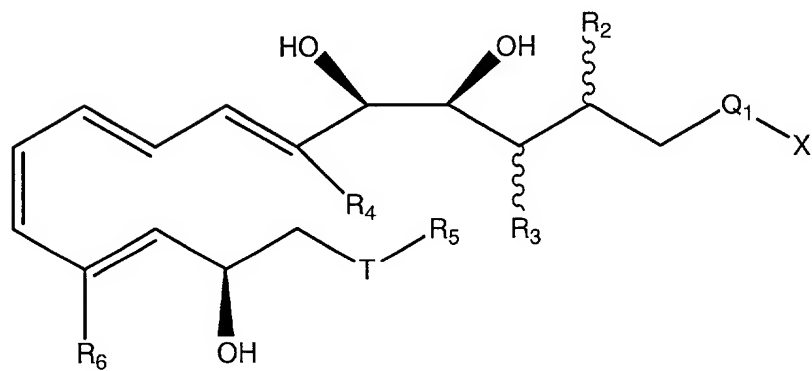


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At page 17, lines 1 through 7, please delete

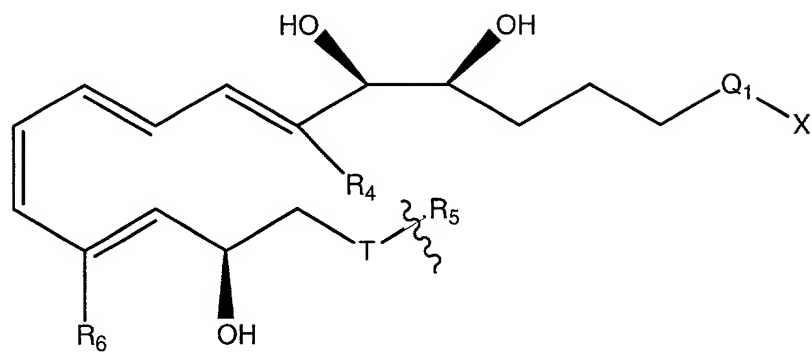


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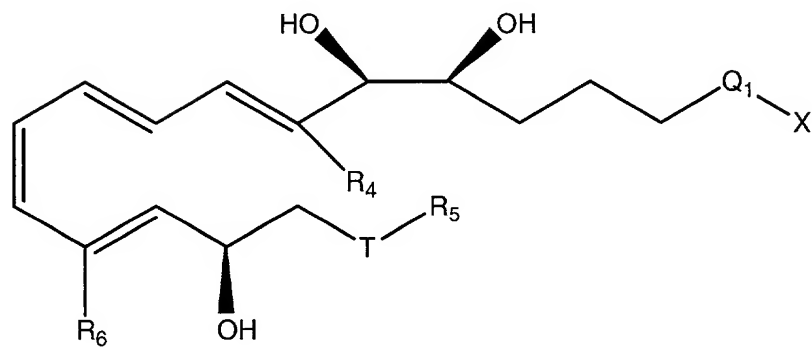


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At page 19, lines 21 through 30, please delete



and insert therefore - -



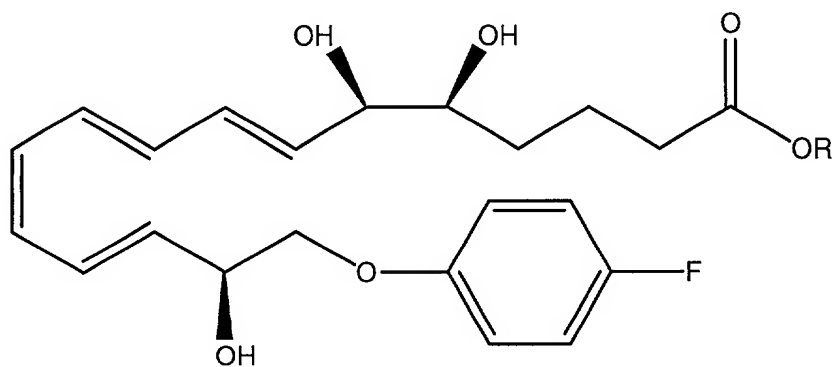
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In the claims:

Please cancel claims 2 through 16 inclusive.

Please add new claims 17 through 32 as follows:

17. (New) A method for modulating a disease or condition associated with phospholipase D (PLD) initiated polymorphoneutrophil (PMN) inflammation in a subject, comprising administering to the subject an effective anti-inflammatory amount of a lipoxin analog having the formula

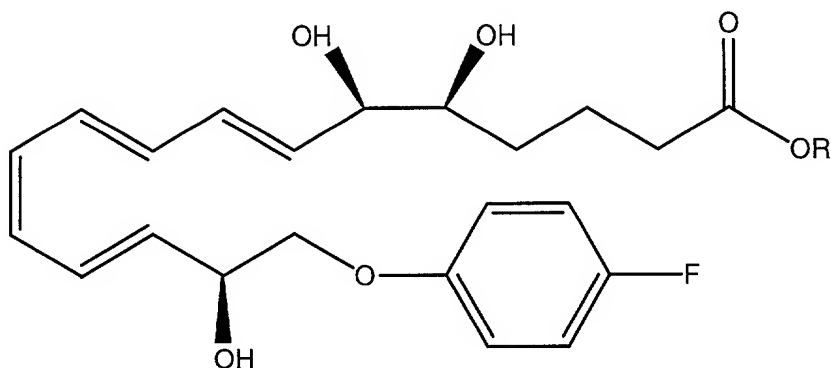


wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof, such that a disease or condition associated with PLD initiated polymorphoneutrophil (PMN) inflammation in a subject is modulated.

18. (New) The method of claim 17, wherein said method is performed *in vitro*.

19. (New) The method of claim 17, wherein said method is performed *in vivo*.

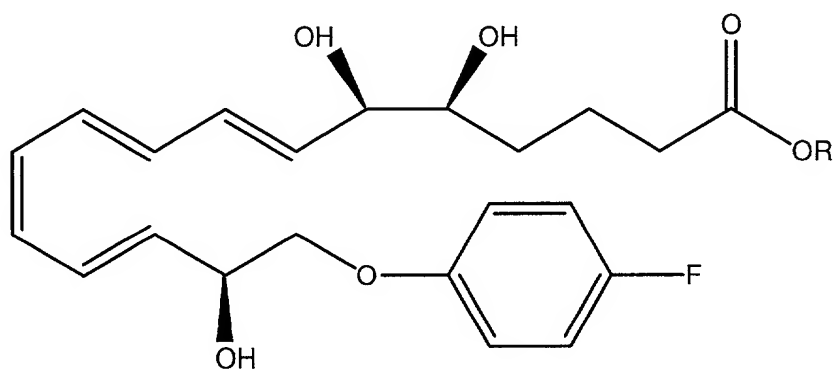
20. (New) A method for treating phospholipase D (PLD) initiated polymorphoneutrophil (PMN) inflammation in a subject, comprising administering to the subject an effective anti-inflammatory amount of a lipoxin analog having the formula



wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof, such that PLD initiated polymorphoneutrophil (PMN) inflammation is treated in a subject.

21. (New) The method of claim 20, wherein said method is performed *in vitro*.
22. (New) The method of claim 20, wherein said method is performed *in vivo*.

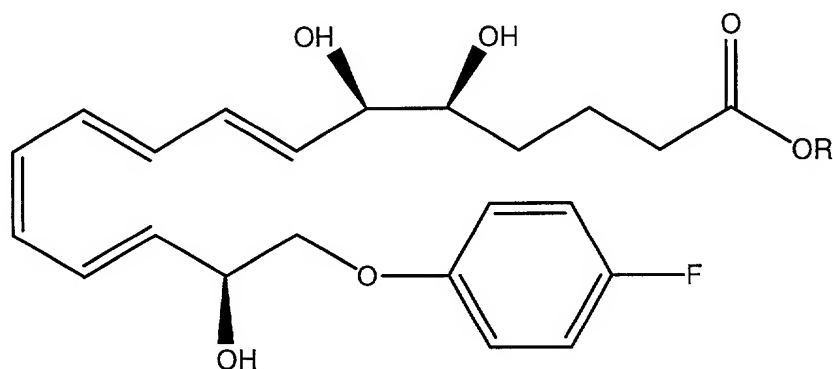
23. (New) A method for modulating a disease or condition associated with phospholipase D (PLD) initiated superoxide generation or degranulation activity in a subject, comprising administering to the subject an effective anti-PLD amount of a lipoxin analog having the formula



wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof, such that a disease or condition associated with PLD initiated superoxide generation or degranulation activity in a subject is modulated.

24. (New) The method of claim 23, wherein said method is performed *in vitro*.
25. (New) The method of claim 23, wherein said method is performed *in vivo*.

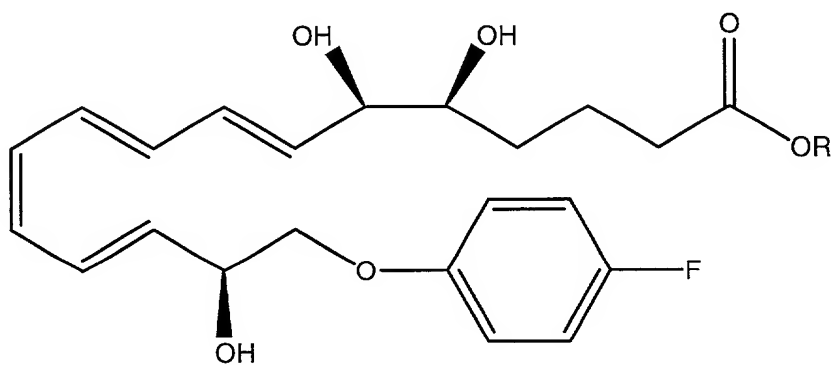
26. (New) A method for treating phospholipase D (PLD) initiated superoxide generation or degranulation in a subject, comprising
administering to the subject an effective anti-PLD amount of a lipoxin analog having the formula



wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof, such that PLD initiated superoxide generation or granulation is treated in a subject.

27. (New) The method of claim 26, wherein said method is performed *in vitro*.
28. (New) The method of claim 26, wherein said method is performed *in vivo*.
29. (New) A packaged pharmaceutical composition for treating a disease or condition associated with phospholipase D (PLD) initiated activity in a subject, comprising:
a container holding a therapeutically effective amount of at least one lipoxin compound having the

formula

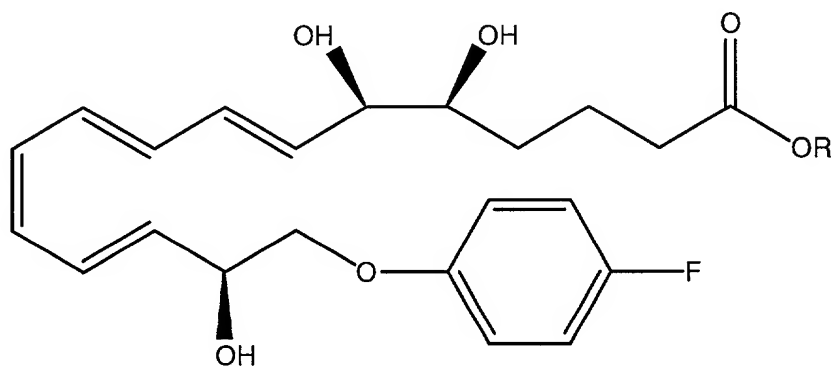


wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating a disease or condition associated with PLD initiated activity in the subject.

30. (New) A packaged pharmaceutical composition for treating phospholipase D initiated activity in a subject, comprising:

a container holding a therapeutically effective amount of at least one lipoxin compound having the formula

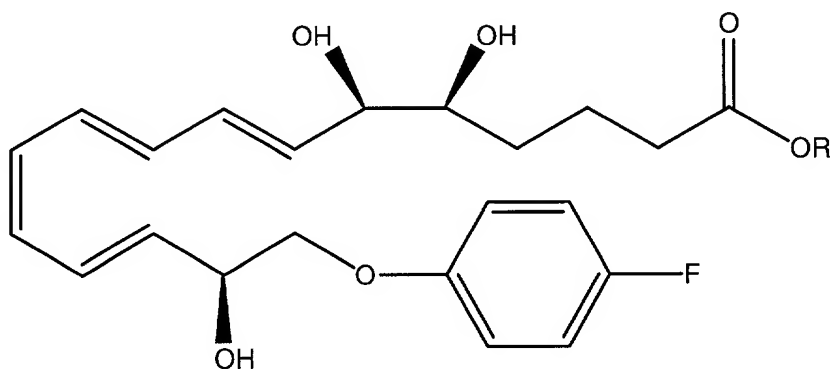


wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating PLD initiated activity in the subject.

31. (New) A packaged pharmaceutical composition for treating a disease or condition associated with phospholipase D (PLD) initiated superoxide generation or degranulation activity in a subject, comprising:

a container holding a therapeutically effective amount of at least one lipoxin compound having the formula

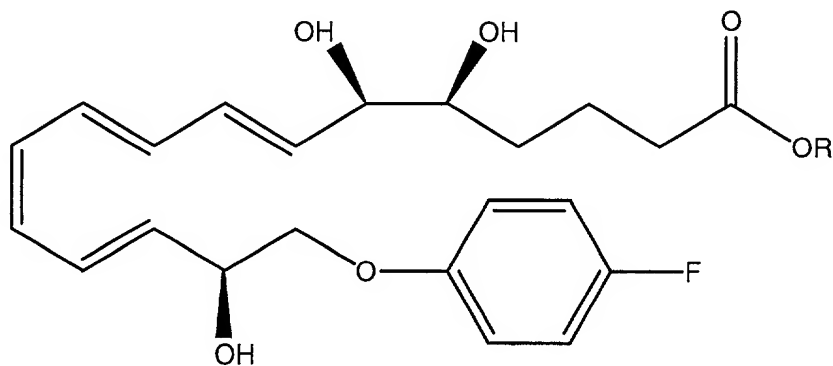


wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating a disease or condition associated with PLD initiated superoxide generation or degranulation activity in the subject.

32. (New) A packaged pharmaceutical composition for treating phospholipase D (PLD) initiated superoxide generation or degranulation activity in a subject, comprising:

a container holding a therapeutically effective amount of at least one lipoxin compound having the formula



wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating PLD initiated superoxide generation or degranulation activity in the subject.

REMARKS

Claims 1 and 17 through 32 are pending.

The specification has been amended to correct for an obvious typographical errors on pages 11, 14, 17 and 19 and to more clearly define the invention.

Attached hereto is a marked up version of the changes made to the claims by the current amendment. The attached pages are captioned "Version with Markings to Show Changes Made."

Conclusion

In view of the foregoing, Applicant submits that all pending claims are allowable. The Examiner is invited to telephone the undersigned attorney for Applicants in the event that such communication is deemed to expedite prosecution of this application.

Respectfully submitted,

DORSEY & WHITNEY LLP

Date: October 19, 2001

By: Scott D. Rothenberger

Scott D. Rothenberger

(Reg. No. 41,277)

DORSEY & WHITNEY LLP

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50 South Sixth Street

Minneapolis, MN 55402-1498

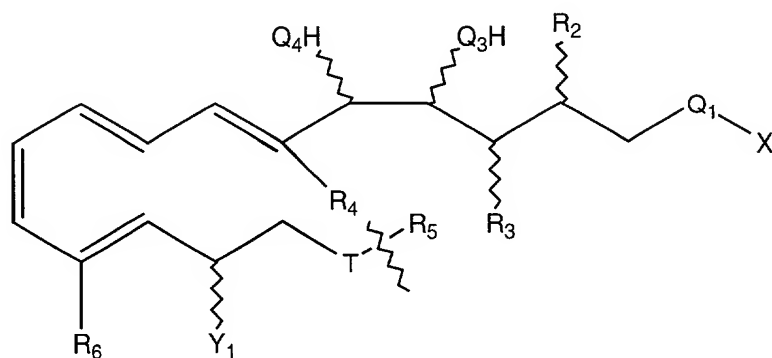
Telephone: (612) 340-8819

Facsimile: (612) 340-8856

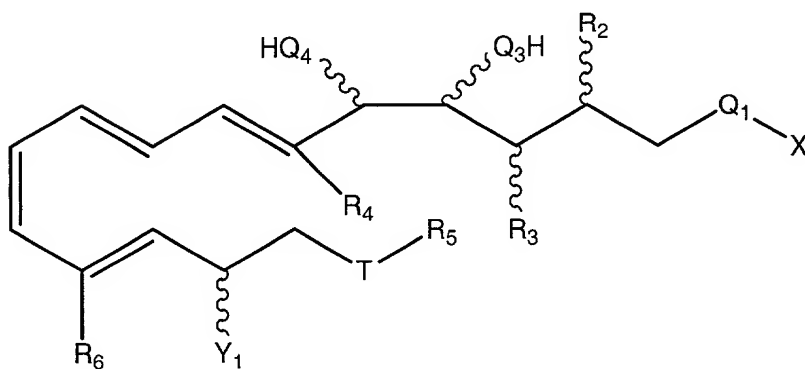
MARKED-UP VERSION SHOWING CHANGES

In the specification:

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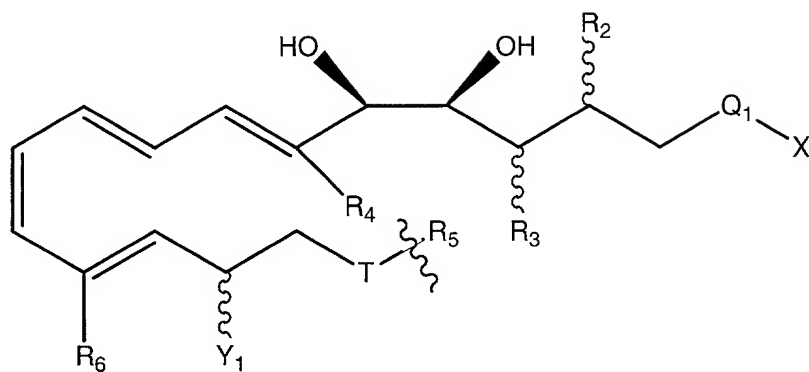


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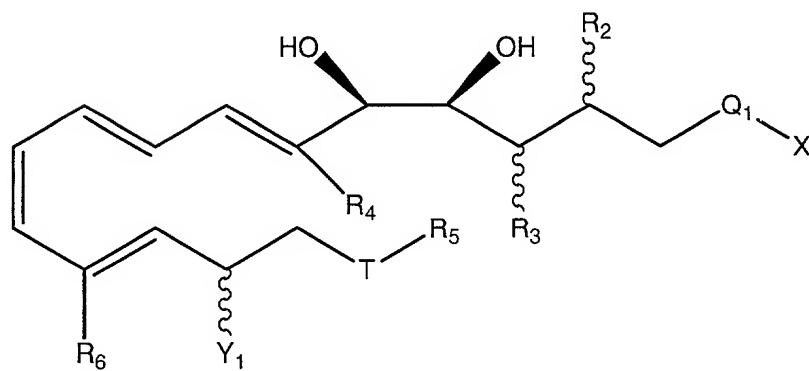


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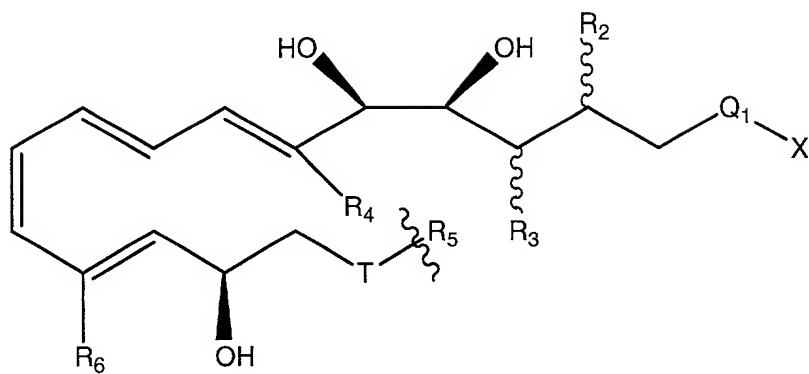


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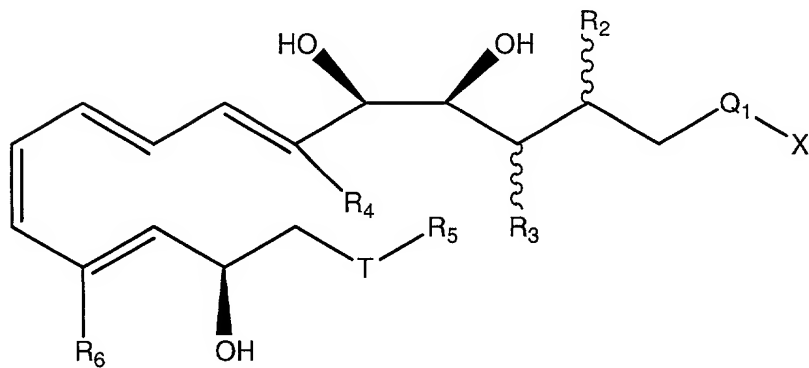


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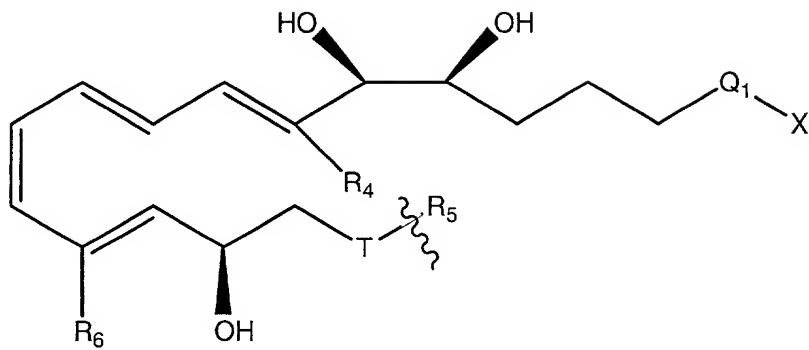


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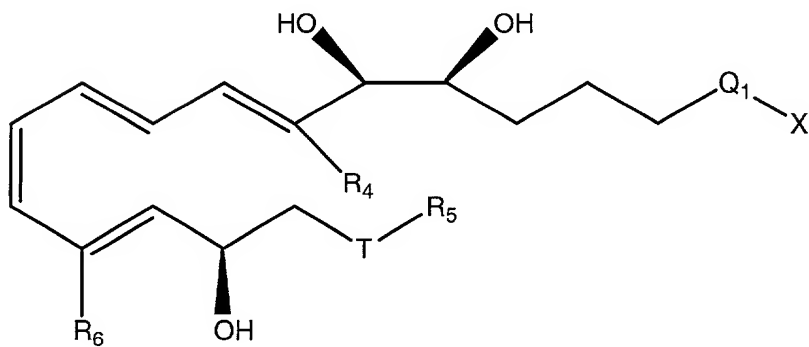


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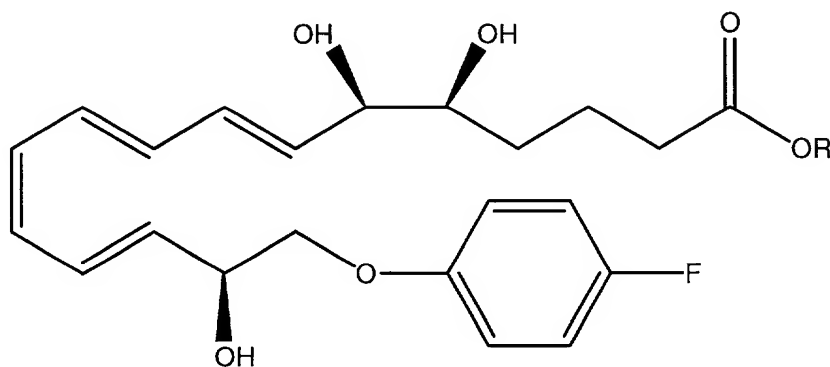
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In the claims:

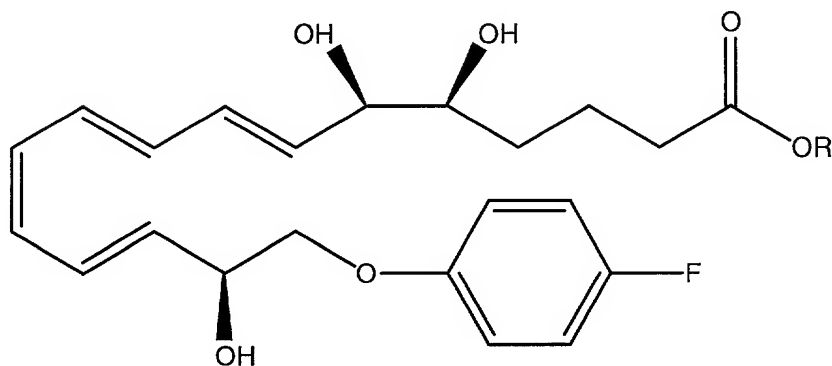
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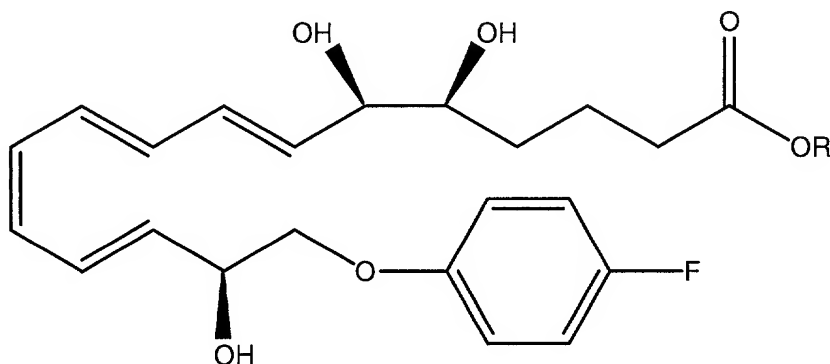
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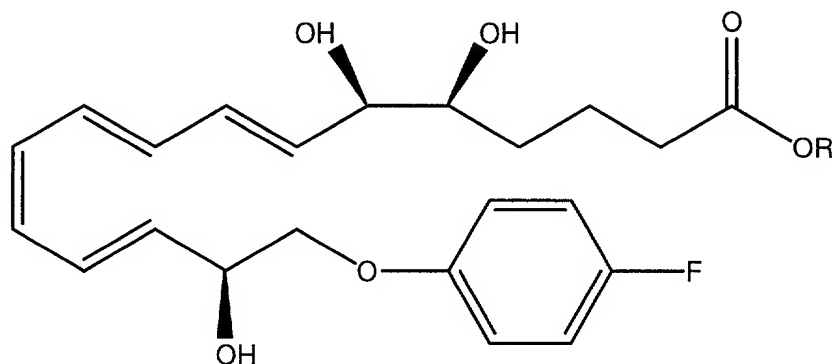
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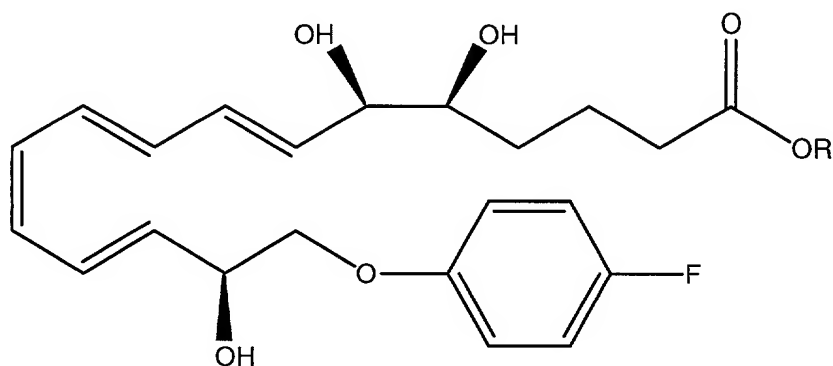
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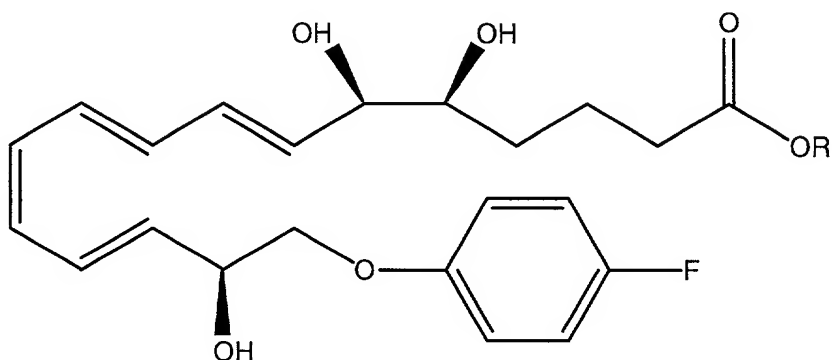


wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating a disease or condition associated with PLD initiated activity in the subject.

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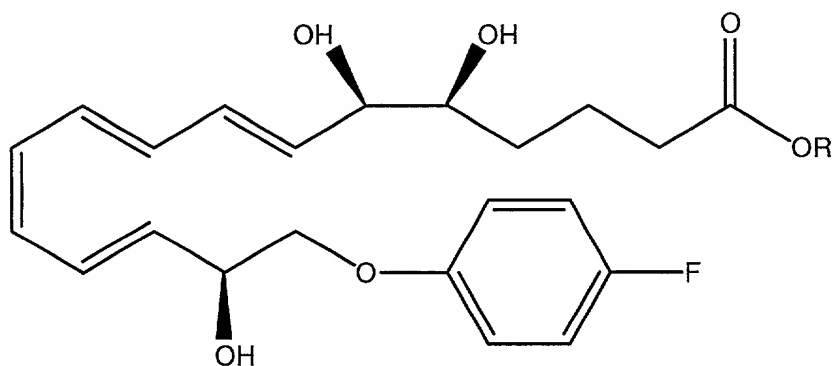


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instructions for using said lipoxin compound for treating PLD initiated activity in the subject.

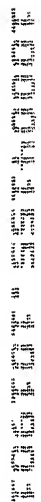
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wherein R is a hydrogen atom, a pharmaceutically acceptable ester and pharmaceutically acceptable salts thereof; and

instructions for using said lipoxin compound for treating a disease or condition associated with PLD initiated superoxide generation or degranulation activity in the subject.

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